



00270

PATENT TRADEMARK OFFICE

Attorney's Docket No.: ACG2AUS

09/700329

TRANSMITTAL LETTER TO THE U.S. ELECTED OFFICE  
(EO/US) - ENTRY INTO NATIONAL STAGE UNDER 35 USC 371

PCT/US99/10751

Int'l Application No.

May 14, 1999

Int'l Filing Date

525 Rec'd PCT/PTO 14 NOV 2000

May 15, 1998

Priority Date Claimed

PREVENTION AND TREATMENT OF HYPERGASTRINEMIA

Title of Invention

Philip C. Gevas, Stephen Grimes, Stephen Karr, Dov Michaeli and Susan Watson

Applicant(s) for EO/US

Box PCT

Assistant Commissioner for Patents

Washington, DC 20231

Attn: EO/US

Sir:

Applicant herewith submits to the United States Elected Office  
(EO/US) the following items under 35 USC 371:

- (1) This express request to immediately begin national examination procedures (35 USC 371(f)).
- (2) A copy of the cover sheet for the published International Application along with a copy of the specification: 42 pages, including 14 claims (2 pages), 12 sheets of formal drawings (Figs. 1-12), 2 pages of Sequence Listing, and a copy of the 4 page International Search Report.
- (3) A copy of the 5 page Request form.
- (4) A Preliminary Amendment.
- (5) An executed Verified Statement Claiming Small Entity Status.
- (6) Our check in the amount of \$345.00, covering the basic national fee as set forth in 37 CFR 1.492(a)(1) (1 total claim; 1 independent; and no multiple dependent).
- (7) A Second Preliminary Amendment and transmittal letter.

Express Mail No.

EK992703206US

- (8) Our check in the amount of \$90.00, covering the extra claim fees after entry of the second Preliminary Amendment (30 total claims; 3 independent; and no multiple dependent).
- (9) an unexecuted Combined Declaration and Power of Attorney form.
- (10) A three (3) page substitute Sequence Listing under PatentIn Version 3.
- (11) A 3.5" computer-readable diskette.
- (12) A 1 page Statement under 37 CFR §1.821(f) and §1.825(a) and (b).

Copies of the following items are also enclosed:

- (13) Copy of 4 page Demand for International Preliminary Examination.
- (14) Copy of 4 page Written Opinion.
- (15) Copy of the 5 page International Preliminary Examination Report.

The Combined Declaration and Power of Attorney form will be filed by the appropriate deadline under 37 CFR §1.495(c)(2) with the surcharge under 37 CFR §1.492(e).

Please charge any additional fees which may be required to effect entry into the National Phase and credit any overpayments to our deposit account 08-3040.

Please direct all communications concerning this application to the undersigned.

Respectfully submitted,  
HOWSON AND HOWSON  
Attorneys for the Applicants

By Mary E. Bak 11/14/00  
Mary E. Bak  
Registration No. 31,215  
Spring House Corporate Center  
Box 457  
Spring House, PA 19477  
Telephone: (215) 540-9206  
Telefacsimile: (215) 540-5818

ACG2AUSA

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re the Application of ) Group Art Unit:  
Philip C. Gevas et al ) Examiner:  
Appln. No. )  
Filed: Herewith )  
For: PREVENTION AND TREATMENT OF ) November 14, 2000  
HYPERGASTRINEMIA )

Assistant Commissioner for Patents  
Washington, DC 20231

**PRELIMINARY AMENDMENT**

Sir:

Please amend the application as set forth below.

In the Claims

Cancel claims 1-6 and 8-14.

REMARKS

After entry of this amendment, the claim pending is claim 7. Claims 1-6 and 8-14 are canceled. No new matter is introduced by this amendment.

Applicants respectfully request that this preliminary amendment be entered prior to calculating the filing fees.

Express Mail No. EK992703204US

The Director of the U. S. Patent and Trademark Office is hereby authorized to charge any deficiency in any fees due with the filing of this paper or credit any overpayment in any fees to our Deposit Account No. 08-3040.

Respectfully submitted,

HOWSON AND HOWSON  
Attorneys for Applicant

By Mary E. Bak  
Mary E. Bak  
Registration No. 31,215  
Spring House Corporate Center  
Box 457  
Spring House, PA 19477  
Telephone: (215) 540-9200  
Telefacsimile: (215) 540-5818

09/700329

529 Rec'd PCT/PTC 14 NOV2000

ACG2AUSA

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re the Application of ) Group Art Unit:  
Philip C. Gevas et al ) Examiner:  
Appln. No. )  
Filed: Herewith )  
For: PREVENTION AND TREATMENT OF ) November 14, 2000  
HYPERGASTRINEMIA )

Assistant Commissioner for Patents  
Washington, DC 20231

**SECOND PRELIMINARY AMENDMENT**

Sir:

Please amend the application as follows.

**In the Specification**

Page 1, line 3, before "Field of the Invention", insert the following:

-- **Cross-Reference to Related Applications**

This is a 371 of PCT/US99/10751, which claims the benefit of the  
priority of US Patent Application No. 60/085,714, filed May 15, 1998 --.

Page 18, line 21, after "-Cys", insert -- [SEQ ID NO: 5] --.

Page 18, line 23, after "-Cys", insert -- [SEQ ID NO: 6] --.

Page 18, line 25, after "-Cys", insert -- [SEQ ID NO: 7] --.

Page 18, line 27, after "-Cys", insert -- [SEQ ID NO: 8] --.

Express Mail No. EK99270320645

Page 18, line 29, after “-Cys”, insert -- [SEQ ID NO: 2] --.  
Page 18, line 31, after “-Cys”, insert -- [SEQ ID NO: 9] --.  
Page 20, line 25, after “-Glu-)”, insert -- [SEQ ID NO: 1] --.  
Page 20, line 26, after “-Cys)”, insert -- [SEQ ID NO: 3] --.  
Page 21, line 4, after “-Glu”, insert -- [SEQ ID NO: 1] --.  
Page 21, line 5, after “-Cys”, insert -- [SEQ ID NO: 2] --.  
Page 21, line 6, after “-Cys)”, insert -- [SEQ ID NO: 3] --.  
Page 21, line 8, after “-Glu”, insert -- [SEQ ID NO: 1] --.  
Page 21, line 11, after “-Glu)”, insert -- [SEQ ID NO: 1] --.

Please enter the attached Abstract of the Disclosure on the attached page.

#### In the Claims

Add new claims 15-43 as follows.

15. A method for treating hypergastrinemia in a mammal comprising administering to said mammal with hypergastrinemia an effective amount of an immunogenic composition that reduces the circulating hormone, gastrin.

16. The method according to claim 15, wherein said immunogenic composition comprises a peptide conjugated to an immunogenic carrier and a pharmaceutically acceptable carrier.

17. The method according to claim 16, wherein said immunogenic composition is selected from the group consisting of a G17 peptide fragment SEQ ID NO: 1 linked by an amino acid spacer to an immunogenic carrier; a G34 peptide fragment SEQ ID NO: 2 linked by an amino acid spacer to an immunogenic carrier; a combination of said G17 and G34 fragments linked by an amino acid spacer to an

immunogenic carrier.

18. The method according to claim 17, wherein said carrier is selected from the group consisting of diphtheria toxoid, tetanus toxoid, and keylimpet hemocyanin.

19. The method according to claim 15, wherein said composition comprises anti-gastrin antibodies that bind to gastrin.

20. The method according to claim 19, wherein said antibodies are purified or humanized.

21. The method according to claim 19, wherein said antibodies bind to heptadecagastrin G17.

22. The method according to claim 19, wherein said antibodies bind to tetratriacontagastrin G34.

23. The method according to claim 19, wherein said antibodies comprise a mixture of antibodies that bind to heptadecagastrin G17 and antibodies that bind to tetratriacontagastrin G34.

24. The method according to claim 15, further comprising administering to said mammal an agent selected from the group consisting of a histamine H<sub>2</sub> receptor blocker and a proton pump inhibitor.

25. The method according to claim 24, wherein said blocker is selected from the group consisting of ranitidine, cimetidine, fomatidine, and nizatidine.

26. The method according to claim 24, wherein said inhibitor is

selected from the group consisting of omeprazole, lansoprazole and pantoprazole.

27. The method according to claim 24, wherein said mammal is administered said immunogenic composition before said agent.

28. The method according to claim 24, wherein said composition and agent are co-administered to said mammal.

29. The method according to claim 15, wherein said hypergastrinemia is associated with a condition selected from the group consisting of pernicious anemia, a gastric tumor, a gastric cancer, and a course of therapy with a substance that results in increased gastrin levels.



30. A method for reducing the side effects of anti-ulcer agents in a mammal comprising administering to a mammal receiving an agent that suppresses gastric acid production or secretion an effective amount of an immunogenic composition that reduces the circulating hormone, gastrin.

31. The method according to claim 30, wherein said immunogenic composition comprises a peptide conjugated to an immunogenic carrier and a pharmaceutically acceptable carrier.

32. The method according to claim 31, wherein said immunogenic composition is selected from the group consisting of a G17 peptide fragment SEQ ID NO: 1 linked by an amino acid spacer to an immunogenic carrier; a G34 peptide fragment SEQ ID NO: 2 linked by an amino acid spacer to an immunogenic carrier; a combination of said G17 and G34 fragments linked by an amino acid spacer to an immunogenic carrier.

33. The method according to claim 32, wherein said carrier is selected from the group consisting of diphtheria toxoid, tetanus toxoid, and keylimpet hemocyanin.

34. The method according to claim 30, wherein said composition comprises anti-gastrin antibodies that bind to gastrin.

35. The method according to claim 34, wherein said antibodies are purified or humanized.

36. The method according to claim 34, wherein said antibodies bind to heptadecagastrin G17.

37. The method according to claim 34, wherein said antibodies bind to tetratriacontagastrin G34.

38. The method according to claim 34, wherein said antibodies comprise a mixture of antibodies that bind to heptadecagastrin G17 and antibodies that bind to tetratriacontagastrin G34.

39. The method according to claim 30, wherein said agent is selected from the group consisting of a histamine H<sub>2</sub> receptor blocker and a proton pump inhibitor.

40. The method according to claim 39, wherein said blocker is selected from the group consisting of ranitidine, cimetidine, famotidine, and nizatidine.

41. The method according to claim 39, wherein said inhibitor is selected from the group consisting of omeprazole, lansoprazole and pantoprazole.

42. The method according to claim 30, wherein said mammal is administered said immunogenic composition before said agent.

43. The method according to claim 30, wherein said composition and agent are co-administered to said mammal.

#### REMARKS

Upon entry of this preliminary amendment, the claims pending are claims 7 and 15-43. New claims 15-43 are supported throughout the specification and by original claims 1-14.

Specification pages 18, 20 and 21 are amended to insert SEQ ID NOS.

The attached Abstract of the Disclosure is supported throughout the specification.

No new matter is introduced by this preliminary amendment and Applicants respectfully request consideration of the pending claims.

A check in the amount of \$90.00 covering the extra claim fees is attached herewith. The Director of the U. S. Patent and Trademark Office is hereby authorized to charge any deficiency in any fees due with the filing of this paper or credit any overpayment in any fees to our Deposit Account No. 08-3040.

Respectfully submitted,

HOWSON AND HOWSON  
Attorneys for Applicant

By Mary E. Bak  
Mary E. Bak  
Registration No. 31,215  
Spring House Corporate Center  
Box 457  
Spring House, PA 19477  
(215) 540-9200

## ABSTRACT OF THE DISCLOSURE

Serum-associated hypergastrinemia is treated by administration of gastrin active or passive immunization. An anti-gastrin immunogenic composition comprising a gastrin G17 or G34 peptide fragment which is amino acid spacer-linked to an immunogenic carrier, is administered so as to effectively neutralize the circulating gastrin hormone, and moreover, inhibit autocrine activity by progastrin such as Gly-extended G17, and amidated G17, in patients with pernicious anemia. Moreover, the method includes administration of a therapeutically effective amount of anti-G17 or anti-G34 antibodies which may be in humanized form. Finally, the method provides ameliorating treatment of hypergastrinemic effects of proton pump inhibitors or H<sub>2</sub> histamine receptor blocking agents or antagonists, in addition to treatment of hypergastrinemia caused by diseases such as pernicious anemia.

Attorney Docket No.: ACG2AUSA

Applicant or Patentee: Philip C. Gevas, Stephen Grimes, Stephen Karr, Dov Michaeli and Susan Watson

Serial or Patent No.: \_\_\_\_\_

Filed or Issued: Filed Herewith

For: PREVENTION AND TREATMENT OF HYPERGASTRINEMIA

VERIFIED STATEMENT (DECLARATION) CLAIMING SMALL ENTITY STATUS  
(37 CFR 1.9(f) and 1.27(e)) - SMALL BUSINESS CONCERN

I hereby declare that I am

- ☐ the owner of the small business concern identified below;  
☒ an official of the small business concern empowered to act on behalf of  
the concern identified below:

NAME OF CONCERN Apton Corporation

ADDRESS OF CONCERN Brickell Bay View Center, 80 S.W. Eighth Street,  
Suite 2160, Miami, Florida 33130-2492

I hereby declare that the above identified small business concern qualifies as a small business concern as defined in 13 CFR 121.3-18, and reproduced in 37 CFR 1.9(d), for purposes of paying reduced fees under Section 41(a) and (b) of Title 35, United States Code, in that the number of employees of the concern, including those of its affiliates, does not exceed 500 persons. For purposes of this statement, (1) the number of employees of the business concern is the average over the previous fiscal year of the concern of the persons employed on a full-time, part-time or temporary basis during each of the pay periods of the fiscal year and (2) concerns are affiliates of each other when either, directly or indirectly, one concern controls or has the power to control the other, or a third party or parties controls or has the power to control both.

I hereby declare that rights under contract or law have been conveyed to and remain with the small business concern identified above with regard to the invention, entitled Prevention and Treatment of Hypergastrinemia by inventor(s) Philip C. Gevas, Stephen Grimes, Stephen Karr, Dov Michaeli, and Susan Watson described in

- ☒ the specification filed herewith  
☐ application serial no. \_\_\_\_\_, filed \_\_\_\_\_  
☐ patent no. \_\_\_\_\_, issued \_\_\_\_\_

If the rights held by the above identified small business concern are not exclusive, each individual, concern or organization having rights to the invention is listed below\* and no rights to the invention are held by any person, other than the inventor, who could not qualify as a small business concern under 37 CFR 1.9(d) or by any concern which would not qualify as a small business concern under 37 CFR 1.9(d) or a nonprofit organization under 37 CFR 1.9(e).

\*NOTE: Separate verified statements are required from each named person, concern or organization having rights to the invention averring to their status as small entities. (37 CFR 1.27)

NAME \_\_\_\_\_

ADDRESS \_\_\_\_\_

☒ INDIVIDUAL ☐ SMALL BUSINESS CONCERN ☐ NONPROFIT ORGANIZATION

I acknowledge the duty to file, in this application or patent, notification of any change in status resulting in loss of entitlement to small entity status prior to paying, or at the time of paying, the earliest of the issue fee or any maintenance fee due after the date on which status as a small entity is no longer appropriate. (37 CFR 1.28(b))

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application, any patent issuing thereon, or any patent to which this verified statement is directed.

NAME OF PERSON SIGNING Philip C. Gevas

TITLE President

ADDRESS OF PERSON SIGNING Apton Corporation, Brickell Bay View Center, 80 S.W. Eighth Street,  
Suite 2160, Miami, Florida 33130-2492

SIGNATURE \_\_\_\_\_

DATE 11-9-00

PREVENTION AND TREATMENT OF HYPERGASTRINEMIAFIELD OF THE INVENTION

The invention relates to the prevention and/or treatment of hypergastrinemia by immunological control of gastrin levels.

BACKGROUND OF THE INVENTION

In humans, treatment with proton pump inhibitors, infection with *Helicobacter pylori* and pernicious anemia account for the majority of cases of hypergastrinemia. Marked hypergastrinemia is seen in the relatively infrequent Zollinger-Ellison Syndrome (ZES). One of the direct effects of hypergastrinemia is, of course, high secretion rates of gastric acid in the stomach.

Around 90% of patients with pernicious anemia (PA) are hypergastrinemic and total gastrin levels can be up to forty times higher than normal levels. A recent study by Varro et al, *J. Clin. Invest.*, 95:1642-1649 (1995) has demonstrated that the hypergastrinemia associated with PA is composed of substantially elevated amidated gastrin, and moderate elevations in the precursors progastrin and glycine extended G17 (gly-G17).

Gastrin peptides are the products of extensive post-translational processing as outlined in Fig. 1. The first translation product of a single mRNA of 0.7 kb is the 101 amino acid precursor preprogastrin. This peptide is translocated into the lumen of the rough endoplasmic reticulum where it is converted into the progastrin peptide. The progastrin moves through the secretory pathway to the golgi stack, and is sulfated at Tyr<sup>87</sup> prior to endoproteolytic cleavage and maturation in the secretory granules. As a consequence, progastrin is processed to give G34 from dibasic cleavage at sites Arg<sup>57</sup> Arg<sup>58</sup> and Arg<sup>94</sup> Arg<sup>95</sup> and to give G17 from dibasic cleavage at sites Lys<sup>74</sup> Lys<sup>75</sup> and Arg<sup>94</sup> Arg<sup>95</sup>. While prehormone convertase 2 (PC2) producing G17 is located primarily in the gastric antrum, the prohormone convertases PC1/PC3 producing G34 are located in the duodenum. The dibasic cleavage residues are removed by carboxypeptidase H (CPH) producing Gly<sup>93</sup> extended gastrins serving as substrates for the amidation enzyme, PAM (peptidylglycine  $\alpha$ -amidating monooxygenase).

Amidated G17 gastrin appears to be a conversion product of G34 NH<sub>4</sub> which is an amidation product of Gly-G34. Gly G17 has been thought as a second endpoint of progastrin processing.

Gastrin effects on tumor cells are via endocrine, paracrine, autocrine and intracrine pathways (Fig. 2) where, however, not all receptor types have been characterized. It is known that exogenous gastrin stimulates gastric and colorectal tumor cells and tumor cell lines.

Most PA patients have endocrine hyperplasia in the gastric corpus and fundus. There is a significant positive correlation between the degree of hypergastrinemia and the number of enterochromaffin-like (ECL) cells. However, the histological type of ECL cell hyperplasia is not dependent on the degree of hypergastrinemia as there is no significant difference in the gastrin levels in patients with linear or nodular hyperplasia. Once diagnosed, despite continuing elevated gastrin levels, the ECL cell hyperplasia appears to remain stable.

The prevalence rate of gastric carcinoid in endoscopically examined PA patients' ranges from 4 to 7%. Patients with carcinoid are diagnosed as having PA 10 years earlier than the average PA patient. This precedes the diagnosis of the carcinoid tumor by a mean of 10 to 12 years. There is no predictive sign for the occurrence of gastric carcinoids in patients with PA, though mean serum gastrin levels are higher in carcinoid compared to ECL hyperplasia [Brinton et al, Brit. J. Cancer, 59:810-813 (1989)]. The stimulus to undergo malignant transformation is thought to be provided by the autoantibodies present. The tumors appear hormonally dependent. Patients who have undergone antrectomy in order to correct hypergastrinemia, have demonstrated disappearance of hyperplastic polyps, carcinoids or agyrophil micronodules diagnosed endoscopically and/or histologically. The demonstration of complete resolution of ECL-cell carcinoids after antrectomy in some patients confirms the potency of hypergastrinemia as a trophic principle for fundic ECL-cells.

In PA, evidence for an effect of the associated hypergastrinemia on other cancers in the gastrointestinal tract comes only from epidemiological studies. Several studies have looked at the incidence of colorectal cancer in PA. A slight increase in the prevalence of colorectal cancer in the first five years after diagnosis of PA has been reported [Talley et al, Annals Int. Medicine, 111:738-742 (1989)].

Studies have also demonstrated an increased prevalence (approximately 7%) of gastric adenocarcinoma in PA. Hypergastrinemia may be responsible for this observed increase. Even though a correlation to serum gastrin levels cannot be found in the majority of patients with gastric cancer, a correlation to chronic atrophic gastritis is always present.

The epidemiological studies have failed to show a consistent increase in the incidence of colorectal cancer in PA. This may be due to the deficient design of the studies. In each analysis the PA patients were compared to unscreened controls from the general population. A proportion of the controls may be expected to be hypergastrinemic due to either *Helicobacter pylori* infection, atrophic gastritis or following administration of a proton pump inhibitor. The apparent failure to show an increase in the incidence of tumors could be explained by the action of gastrin – it acts as a mitogen not a mutagen. However, as gastrin promotes the proliferation of the normal colonic mucosa, there may be an increased chance of a spontaneous mutation, which would affect tumor incidence.

A single study performed recently has looked at the proliferation rate of cells of the normal colon in patients with PA compared to normal controls [Talley et al, cited above]. The control patients were normogastrinemic and had no colonic abnormalities assessed by colonoscopy or barium enema. Using 5'-bromodeoxyuridine to provide a proliferation index, the percentage of proliferating cells in the entire crypts was similar in both groups. In the PA group there was a significantly higher labeling frequency in the upper two fifths of the glands ( $p < 0.01$ ). Movement of the proliferative compartment is seen in individuals at high risk of cancer.

Long-term treatment with omeprazole is known to induce ECL cell hyperplasia which is related to the serum gastrin level. Chronic hypergastrinemia-related carcinoid tumors of the stomach have been reported in certain animals test subjects, e.g. rats, although not yet confirmed in the human [Sobhani et al, Gastroenterology, 105:22-30 (1993)].

Proton pump inhibitors cause a twofold to fourfold increase in fasting and postprandial plasma gastrin concentrations. The increase in fasting hypergastrinemia occurs within a few months of starting therapy. Occasionally, markedly elevated gastrin levels (10 fold) may develop during long term treatment with omeprazole, e.g.,



20-60mg/day. Gastrin levels stabilize after a few months of therapy even if the dose of omeprazole is decreased from 40mg to 20 mg daily [Sontag et al, Gastroenterology, 102:109-118 (1992)].

The growth of gastric endocrine cells has been extensively monitored in patients treated with 20 to 40mg omeprazole daily for up to eight years. No significant quantitative changes of the antral G- and D- cells have been found even after years of high-dose omeprazole treatment. In comparison, the G-cell volume in rats doubled both qualitatively and quantitatively after four weeks of treatment with omeprazole [Tielemans et al, Gastroenterology, 96:723-729 (1989)]. Only patients with the highest serum gastrin levels (>240 pg/ml, four times the upper limit of normal) showed an increase in gastric ECL-cell volume density between the third and fifth year of therapy. This data supports earlier findings that an increase of the ECL-cell volume density is correlated to elevated fasting serum gastrin levels. In addition, linear and nodular hyperplasia was confined to the group of patients with the highest serum gastrin levels. Dysplasia was not been seen in any patient.

A correlation between different grades of atrophy of the oxyntic mucosa and ECL cell growth has been established. In patients receiving 40mg of omeprazole daily for eight years, it was found that the prevalence of micronodular hyperplasia in superficial corpus gastritis was low, e.g., 3.6%, increasing to 19.6% in interstitial gastritis and to 48% in atrophic gastritis. This relationship between atrophic gastritis and micronodular hyperplasia may partially be explained by condensation of the endocrine cells caused by atrophy of the gastric glands and thus may not represent true hyperplasia. Therefore, excessive long-lasting hypergastrinemia induced by omeprazole leads to only linear and simple hyperplasia. In patients with atrophic gastritis or those with a genetic predisposition, hypergastrinemia gives rise to micronodular hyperplasia under the chronic treatment.

ECL hyperplasia in animal models occurs following the administration of omeprazole. The relative growth of both exocrine and endocrine cells produced by hypergastrinemia varies between species. For example, administration of omeprazole, (400µmol/kg, 14mg/kg) to mice for 10 weeks resulted in a threefold increase in plasma gastrin during treatment. Furthermore, the stomach weight increased by 34% and the ECL density by 37% at the end of treatment. The same dose has been found in rats to

increase the gastrin levels 10-fold, resulting in the same general trophic effect (increase of stomach and mucosal weight) as in mice, but the ECL cell density increases by about 300%. The significance of this imbalance in the trophic effect of gastrin on the exocrine cells and ECL cells for the development of carcinoids in rats is not known.

5           Several studies have investigated the effect of hypergastrinemia on normal colonic epithelial cells. The majority of these studies have induced hypergastrinemia by omeprazole administration or as a result of antral exclusion and have produced conflicting results. The effect of long-term (1-year) treatment of female rats with high dose daily omeprazole (400 $\mu$ mol/kg, 14mg/kg) which led to 15 fold increase in gastrin compared to controls was examined [Sundler et al, in Proc. The First International Symposium on Omeprazole, K. O. Borg et al eds, AB Hässle (1986)]. The mucosal thickness of the colonic mucosa and the number of chromogranin-A-containing endocrine cells were unaffected by the omeprazole-induced hypergastrinemia. However, the same animals developed a modest and stable antral gastrin cell hyperplasia. Similarly, Oscarson demonstrated that long-term changes in endogenous gastrin concentration produced by fundectomy (resulting in a 3.5 fold gastrin elevation) did not result in colonic mucosal trophic effects.

10           In contrast, significantly enhanced proliferation of colonic mucosa in omeprazole treated rats compared to controls was demonstrated [Pawlikowski et al, Hormone & Metab. Res., 21:89-91 (1989)]. Short term hypergastrinemia induced colonic mucosal proliferation as well as chronic endogenous hypergastrinemia were demonstrated in rats [McGregor et al, Annals Surg., 195:219-223 (1982)]. Chronic hypergastrinemia was achieved by antral exclusion and short term hypergastrinemia was achieved by pentagastrin administration (2mg/kg) every 12 hours for 48 hours prior to sacrifice. Tissue content and synthesis of DNA, RNA, and protein were all markedly increased by both endogenous gastrin and exogenous pentagastrin. The stimulation by gastrin was significantly stronger than that of pentagastrin. Using the metaphase-arrest technique it has also been have shown that an enhanced mitotic activity of the colonic mucosal cells in rats treated with omeprazole compared to controls [Lewinski et al].

15           More compelling evidence for a trophic role of gastrin has been provided by the development of gastrin deficient transgenic mice. These mice are incapable of

producing gastrin mRNA and the gastrin peptide. This deficiency has allowed studies on the effect of gastrin on the growth and development of the gastrointestinal tract.

Combining histology and immunohistochemical techniques, together with bromodeoxyuridine incorporation, the effect of exogenous gastrin on colonic architecture was assessed. The gastrin deficient mice had histologically normal colons. A decreased proliferation labeling index ( $2.97\% \pm 0.52\%$ ) was noted in such mice compared with wild-type animals ( $4.71\% \pm 0.44\%$ ;  $P < 0.01$ ). The conclusion from these observations is that gastrin is trophic for the normal colonic mucosa.

According to Tang et al. (1996) carcinoid tumors from the *Mastomys* rodent during progression lose response to exogenous hypergastrinemia but have up-regulated expression of TGF $\beta$ . As TGF $\beta$  autocrine pathway potentially acts in a co-operative way with the gastrin autocrine pathway [Howell et al, (1997)], the lack of response of the carcinoids to exogenous gastrin may reflect the increasing activity of the gastrin autocrine pathway. The gastrin gene is apparently activated to rather a lower extent in adenomas than adenocarcinomas.

The conflicting results produced by the above studies may in part be explained by Wang et al, *J. Clin. Invest.*, 98:1918-1929 (1996), in which evidence was provided that progastrin, once thought to be an inert precursor, also has a trophic effect on colonic mucosa. The study included the use of transgenic mice containing a human gastrin (hGAS) minigene, that expresses abundant human gastrin mRNA and human progastrin in the liver. The hepatocytes are unable to process this peptide to the mature amidated form, resulting in markedly elevated serum progastrin levels and normal amidated gastrin levels. The result was a marked increase in the bromodeoxyuridine labeling index of the colon, but not the gastric mucosa, in hGAS mice compared to age-matched, wild-type control mice. This study suggests that progastrin may contribute to colonic mucosal proliferation *in vivo*. Therefore, in conditions of hypergastrinemia not only may the degree of hypergastrinemia be important but the particular gastrin peptide which is elevated may also play a significant role. Normal colonic epithelial cells do not express classical gastrin/CCKB receptors so the action of gastrin must be mediated by an uncharacterized receptor that mediates the action of gastrin precursors.

As stated above, gastrin acts as a mitogen, and thus would not be expected to cause a cell to mutate. This hypothesis which has been confirmed in transgenic hGAS mouse studies. However, if the mucosa has an enhanced proliferation rate, there may be an increased chance of sporadic mutation. The only example of malignant change in animal models occurring in the presence of hypergastrinemia is carcinoid in rats following long term omeprazole administration. Although this finding is particular to rats, and no other animal model produces spontaneous carcinoids, it was felt that omeprazole may have a direct carcinogenic effect. However, the proton pump inhibitor class of drugs that produce hypergastrinemia, ECL hyperplasia and ECL carcinoids in the rat, has tested negatively for genotoxicity. Subsequent studies have shown that it is not a specific drug that leads to carcinoid formation; carcinoids can also be produced by feeding with 2000mg/kg ranitidine, loxitidine, the hypolipidemic agent clofibrate and by 75% corpectomy, all of which produce hypergastrinemia. The mediator role of gastrin was confirmed when it was shown that antrectomy in rats prevents omeprazole induced ECL cell hyperplasia. The formation of carcinoids in rats simply in the presence of hypergastrinemia may be due to their genetic background.

There is no reported evidence of hypergastrinemia producing spontaneous tumors at other sites in the gastrointestinal tract. In humans, it is evident that an additional factor may be required for ECL cells to progress from simple hyperplasia to carcinoid. In PA, the additional factor is possibly supplied by the presence of autoantibodies.

Once the cell has been transformed, exogenous gastrin can continue to promote growth. This effect may be enhanced by gastrin/CCKB receptors which are expressed de novo on adenomas. The exact point in the adenoma-carcinoma transformation sequence at which the gastrin/CCKB receptor and autocrine gastrin are expressed is not yet known. Hypergastrinemia may increase this transforming progression through the stages of the adenoma-carcinoma sequence.

In addition, treatment with agents directed against excess production of gastric acid has been found to induce parietal cell hyperplasia and hypertrophy. Recent cases were reported to suggest a correlation between gastric acid-inhibitory treatment by either proton pump inhibitors, such as omeprazole, lansoprazole, or histamine H<sub>2</sub> receptor inhibiting agents, such as ranitidine or cimetidine, and the occurrence of

fundic gland polyps (FGP).

A therapeutic method for selectively immunologically neutralizing the biological activity of the gastrin hormone would provide an effective means to control or prevent the physiopathological changes resulting from hypergastrinemia.

5 As disclosed in co-assigned U.S. Patents Nos. 5,609,870; 5,607,676; 5,622,702; 5,468,494; and 5,023,077, immunization against the G17 and G34 gastrin forms can effect neutralization of serum gastrin. The immunogenic constructs of this invention include an aminoterminal (1-9) G17 peptide or an aminoterminal (1-6) G34 peptide conjugated via a peptide spacer to an immunogenic carrier. The preferred G17  
10 sequence is pyro-Glu-Gly-Pro-Trp-Leu-Glu-Glu-Glu-Glu [SEQ ID NO: 1] and the preferred G34 sequence is pGlu-Leu-Gly-Pro-Gln-Gly-Arg-Pro-Pro-Pro-Cys [SEQ ID NO: 2]. The preferred spacer in both constructs is a Ser-peptide (Ser-Ser-Pro-Pro-Pro-Pro-Cys [SEQ ID NO: 3]). The preferred immunogenic carrier is diphtheria toxoid, tetanus toxoid, keylimpet hemocyanin, and bovine serum albumin  
15 (BSA). The gastrin immunogen is defined as a conjugate of the pGlu- Gly-Pro-Trp-Leu-Glu-Glu-Glu-Glu [SEQ ID NO: 1] peptide sequence, with an amino acid spacer linked to an immunogenic carrier. The preferred gastrin immunogen is defined as a conjugate of the (1-9) amino terminal (pGlu-Gly-Pro-Trp-Leu-Glu-Glu-Glu-Glu [SEQ ID NO: 1]) peptide which is linked by peptide spacer to diphtheria toxoid. It is further  
20 known that the gastrin immunogen preparation is also effective for inhibiting the incompletely processed or progastrin type gastrin precursors which may be bound to the cell membrane of a gastrin producing cell.

There is a need in the art for compositions and methods to effectively treat hypergastrinemia.

## 25 SUMMARY OF THE INVENTION

The present invention is directed to the treatment for control or prevention of gastrointestinal disorders such as hypergastrinemia by administering a gastrin immunogen preparation to an afflicted mammal or human.

A preferred embodiment of the treatment is directed to the control or  
30 prevention of hypergastrinemia due to pernicious anemia.

Another preferred embodiment of this invention is directed to the treatment for control or prevention of gastrointestinal side effects due to antiulcer agents such as proton pump inhibitors or histamine H<sub>2</sub> receptor blocking agents or antagonists.

It is another preferred embodiment of this invention to treat hypergastrinemia related to colorectal disorders or diseases by immunization with gastrin immunogen against gastrin peptide G17, G34, amidated gastrin and progastrin. In this context, the anti-G17 immunogen as described in U. S. Patent Nos. 5,609,870; 5,468,494; 5,785,970 and in the co-assigned patent application 08/798,423 has been found to provide an effective agent to stimulate anti-G17 antibodies which cross-react with Gly extended G17 (G17-Gly), amidated G17 (G17 NH<sub>2</sub>) so as to be suitable for treating gastrointestinal tumors which are responsive to these gastrin peptides. The '423 application is incorporated herewith by reference in its entirety.

It is a special advantage of the present invention to provide a specific immunogen or antibody to target the specific protein which results in hypergastrinemia. For example, Gly G17 and G17 NH<sub>2</sub> can be neutralized with an anti-G17 immunogen composition, such as G17 (1-9) Ser DT, while G34 can be neutralized with anti-G34 (1-17) immunogens.

Moreover, G17 and G34 can be neutralized by anti-G34 (13-22) and anti-G34 (17-31) immunogens which generate antibodies able to cross-react with both gastrin epitopes. Passive immunization can be effected by the specific antibodies generated by immunogens against the various G17 and G34 epitope. These antibodies will either react specifically and separately with the G17 or G34 epitopes or react with both such gastrin epitopes together.

It is an especially preferred embodiment of this invention to treat or pre-treat with gastrin immunogen-type immunization a patient or mammal who is under chronic or long term treatment with the proton pump inhibitor, omeprazole or lansoprazole. A further embodiment provides passive immunization with anti-G17 antibodies which may be humanized to treat hypergastrinemia. A perfected combination treatment of hypergastrinemia and concomitant excess product of gastric acid involves administration of proton inhibitors or H<sub>2</sub> histamine receptor blockers.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 illustrates the processing of the gastrin precursor to the mature gastrin forms;

Fig. 2 illustrates the various pathways of gastrin activity;

5 Fig. 3 illustrates the structural aspects of a gastrin immunogen.

Fig. 4 illustrates bound gastrin (median) in normal and hypergastrinemic subjects (the immunogen control group did not have bound serum gastrin above the 10 pg/ml detection limit).

10 Fig. 5 illustrates percentage survival of experimental hypergastrinemia mice treated with gastrin immunogen compared to controls.

Fig. 6 depicts the mean proliferation index of gastrin associated tumors in Min mice under treatment with gastrin immunogen.

Fig. 7 illustrates the timed levels of antibody in Min mice immunized with G17 (1-9):DT.

15 Fig. 8 compares the Min mouse anti-G17 (1-9):DT antibody levels in response to hG17-DT immunogen + vehicle, hG17-DT immunogen + omeprazole, vehicle only, omeprazole only, positive and negative controls.

20 Fig. 9 compares Min mouse serum G17 levels when immunized with 1) hG17-DT immunogen plus vehicle (Free G17), 2) plus vehicle (Bound G17), 3) plus omeprazole (free G17) and 4) plus omeprazole (Bound G17).

Fig. 10 illustrates the percent animals surviving after treatment with oral vehicle plus blank immunogen (n=22); omeprazole (n=18) plus blank immunogen (n=30), oral vehicle plus gastrin immunogen (n=18); and omeprazole hG17-DT immunogen (n=30).

25 Fig. 11 shows the displacement of labelled G17 from anti-N-terminal gastrin (from rabbit anti-human G7 antiserum) by G17, Gly-G17, and G34 as described in Example 5.

Fig. 12 shows the displacement of labelled G17 from anti-C-terminal gastrin (from rabbit anti-human G7 antiserum) by G17, Gly-G17, and G34 as described in Example 5.

DETAILED DESCRIPTION OF THE INVENTION

As described above, immunogenic constructs useful in this invention include an aminoterminal (1-9) G17 peptide or an aminoterminal (1-6) G34 peptide conjugated via a peptide spacer to an immunogenic carrier. The preferred G17 sequence is pyro-Glu-Gly-Pro-Trp-Leu-Glu-Glu-Glu-Glu [SEQ ID NO: 1] and the preferred G34 sequence is pGlu-Leu-Gly-Pro-Gln-Gly-Arg-Pro-Pro-Pro-Cys [SEQ ID NO: 2]. The preferred spacer in both constructs is a Ser-peptide (Ser-Ser-Pro-Pro-Pro-Pro-Cys [SEQ ID NO: 3]). The preferred immunogenic carrier is diphtheria toxoid, tetanus toxoid, keylimpet hemocyanin, and bovine serum albumin (BSA). The gastrin immunogen is defined as a conjugate of the pGlu- Gly-Pro-Trp-Leu-Glu-Glu-Glu-Glu [SEQ ID NO: 1] peptide sequence, with an amino acid spacer linked to an immunogenic carrier. The preferred gastrin immunogen is defined as a conjugate of the (1-9) amino terminal (pGlu-Gly-Pro-Trp-Leu-Glu-Glu-Glu-Glu [SEQ ID NO: 1]) peptide which is linked by peptide spacer to diphtheria toxoid.

In addition to the above-named anti-gastrin immunogens, one of the generating antibodies binding to both G17 and G34 comprises a conjugate of the 7 amino acids of C-terminal G17 amino acid sequence 11-17-DT. This sequence is E-A-Y-G-W-M-D-NH<sub>2</sub> [SEQ ID NO: 4]. An inhibition of G34 and G17 induced gastric acid production in perfused rat stomachs was observed after an intravenous injection of 1ml of rabbit Anti-C terminal G17 (11-17)-DT antisera.

In order to explore whether gastrin may promote progression to malignancy in existing pre-malignant conditions, studies were undertaken, for example, in the multiple intestinal neoplasia or so-called *Min* mouse model of familial adenomatous polyposis (FAP). The mice have a germline mutation in their APC gene which leads to multiple intestinal neoplasia. Hypergastrinemia which was induced by use of the proton pump inhibitor, omeprazole, has been now found to increase progression to malignancy in *Min* mice, reducing their median survival from approximately 10 weeks to 6 weeks. Examination of the proliferation of tumors from *Min* mice exposed to elevated gastrin levels revealed by bromodeoxyuridine incorporation that proliferation was increased.



In addition to the proliferative effects of serum-associated gastrin, acting in an endocrine manner to increase proliferation, expression of the gastrin gene has also been shown in the colonic mucosa in pre-malignant condition. In the transgenic APC1628 mouse, the gastrin gene is activated in both the normal colonic mucosa and the malignant epithelium [Smith et al, Brit. J.Surg., 84:706 (1997)]. This has recently been confirmed by applicants by both immunocytochemistry and at the gene level in the *Min* mouse. In addition, activation of the gastrin gene has been found in human adenomas [Smith et al, (1997) cited above]. Thus, a gastrin-mediated autocrine/paracrine pathway may also be operational in the pre-malignant scenario.

Effect of gastrin neutralization on the progression of the adenoma:carcinoma sequence in the *Min* mouse model of familial adenomatous polyposis has been observed. Specifically, both serum-associated gastrin and gastrin present with the colonic epithelium may contribute towards the progression cascade in the *Min* mouse model of FAP. Accordingly, the effect of gastrin neutralization on *Min* mouse survival was determined.

The *Min* mice used in the study were bred within the Academic Unit of Cancer Studies at Nottingham University (U.K.) on a C57/BL background. As the homozygous state is lethal and female *Min* mice do not lactate, a *Min* heterozygote is bred with a female wild type and 1:4 offspring have the *Min* genotype. The *Min* positive mice were then placed into each arm of the therapy on an ongoing basis.

The immunization with hG17-DT immunogen (G-17 conjugate) (Fig. 3), is effective in neutralizing circulatory gastrin levels as well as tissue bound precursors or incompletely processed progastrin.

Using exogenous anti-G17 antibodies which can be in humanized form, a patient can be preimmunized against hypergastrinemia or hypergastrinemic effects caused by treatment with proton inhibitors (omeprazole, lansoprazole, or pantoprazole) or H<sub>2</sub> receptor blockers (ranitidine, cimetidine, formatidine or nizatidine). Humanized antibodies may be prepared by techniques known in the art.

The hG17-DT conjugated immunogen or method of preparation are disclosed in U.S. Patent No. 5,609,870, U.S. Patent No. 5,468,494 and U.S. Patent No. 5,023,077 which are incorporated by reference in this specification in their entirety. See, also, Examples 3 and 4 below.

The endpoint of the treatment, which has previously been validated by Moser et al. (1990), is at the terminal stage of the disease, when the mice have a large tumor burden, blood is lost in the stools, and the animals become anemic.

The following examples are provided for illustration only.

5      Example 1

Gastrin neutralization was achieved by using an immunogen, i.e. the gastrin immunogen preparation, which is composed of the amino terminal domain of gastrin-17 linked, via an amino acid spacer, to diphtheria toxoid which acts as the immunogenic carrier. The antibodies raised by virtue of the design of the immunogen, cross-react with both amidated and glycine-extended gastrin-17, known proliferative forms of gastrin. *Min* mice were immunized s.c. with the hG17-DT immunogen (100 mg/mouse) at week 4, with subsequent injections at 3 weekly intervals. Serum antibody titres are known to rise within 2 weeks of the first immunization at levels with an antigen binding capacity of  $> 10^{-9}$ M. The hG17-DT immunogen was administered to mice at 4 weeks of age to examine its effect on mice with an established tumor burden. Control mice received immunogen constituents without the active peptide.

The presence of anti-gastrin antibodies within the serum of gastrin immunogen-immunized mice was confirmed by using an ELISA capture assay. To confirm the presence of antibody-bound gastrin, serum was taken from immunized mice, antibody:antigen complexes were purified, uncomplexed by boiling and the bound gastrin measured by RIA.

Bound gastrin levels were not measurable in animals immunized with control immunogen. The bound gastrin levels in the gastrin immunogen-immunized mice were 37pg/ml. In hypergastrinemic mice, with a 3-4 fold increase in serum gastrin levels, the bound gastrin levels was 148.3pg/ml which highlights the capacity of the antibodies raised in the serum for gastrin neutralization (Fig. 5).

There was a significant effect on survival, with the median survival in the control immunogen treated group being 8 weeks compared to 15 for the gastrin immunogen treated mice (Fig. 6). Log rank test  $p=0.0017$ . However, there is a sharp decline in the survival of the mice after 14-15 weeks.

One hour prior to termination, mice were injected with DNA analogue bromodeoxyuridine in order to determine the comparative in situ proliferation activity. The samples were formalin fixed, paraffin embedded, sections were cut and stained with an anti-bromodeoxyuridine antibody. There was no significant difference in the proliferation of crypts from both the small and large intestine due to gastrin immunogen treatment (Fig. 7). The proliferation of the small intestinal tumors were significantly inhibited by 19.8%, as was the proliferation of the large intestinal tumors which was inhibited by 41%.

The normal colonic mucosa is sensitive to the proliferative effects of hypergastrinemia involving both amidated gastrin and progastrin (Wang, et al., 1997; Renga et al., 1997). APC1638 and *Min* mice (both have mutations in their APC gene) have an activated gastrin gene in both normal and malignant colonic mucosa, unlike the corresponding wild type mice. *Min* mice have greater proliferation levels in normal mucosa when compared to the wild type C57/BL mouse.

Hypergastrinemia induced by treatment with high daily doses of omeprazole decreases the survival of *Min* mice, which is partially reversed when co-administered with gastrin immunogen. There is an initial 2 week window when hypergastrinemia is unopposed due to lack of anti-gastrin antibodies. This effect is completely reversed when omeprazole treatment is delayed for 2 weeks allowing anti-gastrin antibody titers to rise prior to onset of hypergastrinemia.

Administration of gastrin immunogen has no effect on survival of mice of increased age (6-12 weeks). However, there is a significant effect on survival of mice immunized at an earlier age (4 weeks). Therefore, the results suggest a stage through which the mice pass and after which an anti-proliferative effect is not enough to suppress malignant progression. The subject mice thus would respond to serum-associated gastrin until onset of the gastrin autocrine pathway at a later age which may be more refractory to gastrin immunogen-induced antibodies.

When gastrin immunogen was given 2 weeks prior to omeprazole treatment a complete reversal of the survival effect of omeprazole-induced hypergastrinemia on *Min* mice was observed. This was confirmed when it was shown that the omeprazole + gastrin immunogen treated group was not significantly different from the vehicle control ( $p=0.1103$ ).

Anti-G17 antibodies had previously been shown to be detectable in 4 week old *Min* mice by week 2 (Fig. 6) following immunization with gastrin immunogen.

Determination of the anti-G17 antibody levels at the termination of the study revealed a variation in both groups as measured by specific absorbance using an ELISA assay.

Some suppression of the levels of anti- G-17 antibodies in mice co-treated with omeprazole was observed perhaps due to the neutralization of the omeprazole-induced hypergastrinemia (Fig. 7,  $p=0.02$ , Mann Whitney, comparing anti-G17 (1-9):DT antibody levels in vehicle and omeprazole-treated groups).

The *Min* mouse anti-G17(1-9):DT antibody data are illustrated in Fig. 5.

Measurement of the free and bound serum carboxy-amidated gastrin levels in immunized animals revealed a mean free gastrin level of 28.9pg/ml and a bound level of 36.7pg/ml (Fig. 8).

In the mice treated with gastrin immunogen + omeprazole, the free gastrin level was 45.6pg/ml (1.5 fold increase compared to animals immunized with gastrin immunogen alone,  $p=0.0135$ , mann Whitney). The bound gastrin level in the gastrin immunogen + omeprazole treated mice was 148.3pg/ml which was significantly increased when compared to animals treated with gastrin immunogen alone (2 animals had bound gastrin levels of greater than 200pg/ml)  $p=0.00001$  when compared to bound gastrin levels in the gastrin immunogen treated group and  $p=0.00001$  when compared to free gastrin levels in the gastrin immunogen + omeprazole group, [Mann Whitney]. The *Min* mouse serum G17 data are illustrated in Fig. 8. The data are shown in Table 1.

Table 1

Group	Mean	SD	Statistics (Mann Whitney)
1.	28.9	12.7	1vs2 $p=0.191$
2.	36.7	15.4	1vs3 $p=0.0135$
3.	45.6	19.2	2vs4 $p=0.00001$
4.	148.3	170.9	3vs4 $p=0.00001$

G17-DT immunogen administered alone induced a significant effect on survival (Fig. 9,  $p=0.0017$ ). The effect on survival was greatest in the initial phase of the experiment with time to 50% survival being 9.5 weeks for the vehicle control and 14.5 weeks for

immunogen treated. G17-DT immunogen only induced this beneficial effect on survival until week 14, following which there was an exponential drop.

### Example 2

As described below, the effect of omeprazole induced hypergastrinemia on the progression of the intestinal neoplasia was further studied in the *Min* (multiple intestinal neoplasia) mouse model of polyposis coli.

Confirmed *Min*<sup>+</sup> genotype mice were randomized into 4 groups:

Group 1. OME 75mg/kg daily oral treatment

Group 2. OME + GSI 100mg oral dose/ mouse day 0 and every 3 weeks

Group 3. Oral vehicle + control immunogen

Group 4. Oral vehicle - control immunogen

Serum gastrin level was measured by RIA. Prior to end of treatment, proliferative index was determined by the bromodeoxyuridine method.

Group 1. 236pg/ml of serum gastrin

Group 4. 67pg/ml of serum gastrin

Group 1 hypergastrinemia significantly decreased survival compared to control ( $p=0.0001$ , log rank test) with mice in control group having a 50% survival of 16 weeks compared to 8 weeks in the omeprazole treated group. HG17-DT immunogen induced formation of serum antibodies with antigen binding capacity of  $> 20\text{mg/ml}$  resulting in effective neutralization of the hypergastrinemic state. Gastrin neutralization resulted in a reversal of the survival disadvantage induced by omeprazole ( $p=0.0017$ ).

The hypergastrinemic mice had enhanced proliferation of normal colonic mucosa. It was found that 9.46% proliferating cells increased to 20.1%,  $p<0.05$ , Mann-Whitney and colonic neoplasia (22.3% increased 35.0%,  $p<0.01$ ). Thus, the level of this experimental hypergastrinemia was in the range attained in the humans on a maintenance dose of omeprazole and resulted in enhanced progression through the adenoma:carcinoma sequence. Moreover, gastrin was confirmed as the mediator inducing a state of hyper-proliferation within both normal and neoplastic colonic epithelium. This data demonstrate the need and effectiveness for controlling hypergastrinemia on pre-malignant colon by gastrin immunogen immunization.

Polypoid carcinomas have been established *in vitro* from the large and small intestine of *Min* mice. Proliferation was assessed by a tetrazolium-based colorimetric ELISA assay. It was found that proliferation of both tumor types was not increased by amidated gastrin, but the large intestinal tumor was modestly stimulated by glycine-extended gastrin.

Gastrin immunogen immunization significantly affects the survival of *Min* mice when administered early in their life span. Moreover, the proliferation index of tumors in the large intestine was more extensively inhibited by the G17-DT immunogen than that of tumors arising in the small intestine. In this context, tumors from the large intestine of *Min* mice appear to be more sensitive to the proliferative effects of GlyG17 than tumors from the small intestine. This could be both serum-associated and tumor-associated GlyG17, the latter being due to activation of the gastrin gene in these tumors.

Immunological inhibition by G17-DT immunogen at the terminal stage of the adenoma:carcinoma sequence was not as effective. As the small intestinal tumors are the most prolific in terms of number and total tumor burden, an inhibitory effect on proliferation of less than 20% discussed above, may not be great enough to stabilize progression.

Thus, it is clear that the above-described results lead to the following conclusions:

1. The MIN mouse over-expresses the APC gene, the mutation apparently responsible for the adenoma formation a pre-cancerous stage.
2. The adenomas are sensitive to gastrin stimulation especially in early stages and/or young mice.
3. Administration of proton pump inhibitors or H<sub>2</sub> blockers as described, causes hypergastrinemia, hyperproliferation of adenous as and consequently shortened survival.
4. Immunization contemporaneous with omeprazole partially reverses the deleterious effect on survival. Immunization with G17-DT immunogen 2 weeks prior to the proton pump inhibitor administration resulted in complete reversal of the deleterious effects of the drug. In this regimen, the rise in antibody titers coincided with the start of the proton pump inhibitor treatment.

Treatment with the G17-DT immunogen as described is useful in reversing hyper-gastrinemic states induced by a variety of conditions, including, PA, H. pylori, atrophic gastritis, total or partial gastrectomy, treatment with proton pump inhibitors or H<sub>2</sub> blockers. The G17-DT immunogen is potentially effective in protecting the subject mammal, including humans, from induction of cancers responsive to gastrin (colon, stomach, pancreas, and liver).

Immunization against gastrin according to the present method of using hG17-DT induces an effective immune response in humans such that it reduces serum gastrin levels in hypergastrinemic patients to normal or lower levels.

Treatment of PA patients exhibiting hypergastrinemia with immunization (active or passive) against gastrin can be applied alone or in combination with a secondary step of anti-gastric acid administration proton pump inhibitors such as omeprazole or lansoprazole, as well as H<sub>2</sub> receptor blocking agents, such as ranitidine, cimetidine, famotidine or nizatidine.

### Example 3

Immunogens capable of inducing specific immune responses to either G17 or to G34 were prepared by standard solid state synthesis methods. Each peptide was characterized as to amino acid content and purity.

Peptides with the following amino acid sequences were synthesized:

Peptide 1 - Human G17(1-6) ("hG17(6)": pGlu-Gly-Pro-Trp-Leu-Glu-Arg-Pro-Pro-Pro-Pro-Cys

Peptide 2 - Human G17(1-5) ("hG17(5)": pGlu-Gly-Pro-Trp-Leu-Arg-Pro-Pro-Pro-Pro-Cys

Peptide 3 - Human G17(1-4) ("hG17(4)": pGlu-Gly-Pro-Trp-Arg-Pro-Pro-Pro-Pro-Cys

Peptide 4 - Rat G17(1-6) ("rG17(6)": pGlu-Arg-Pro-Pro-Leu-Glu-Arg-Pro-Pro-Pro-Pro-Cys

Peptide 5 - Human G34(1-6) ("hG34(6)": pGlu-Leu-Gly-Pro-Gln-Gly-Arg-Pro-Pro-Pro-Pro-Cys

Peptide 6 - Human G34(13-22) ("hG34/G17 combination"): Asp-Pro-Ser-Lys-Lys-Gln-Gly-Pro-Trp-Leu-Pro-Pro-Pro-Pro-Cys

Each of these peptides were conjugated to amino groups present on a carrier such as Diphtheria toxoid ("DT") via the terminal peptide cysteine residue utilizing heterobifunctional linking agents containing a succinimidyl ester at one end and maleimide at the other end of the linking agent.

5 To accomplish the linkage between any of Peptides 1-6 above and the carrier, the dry peptide was dissolved in 0.1 M Sodium Phosphate Buffer, pH 8.0, with a thirty molar excess of dithiothreitol ("DTT"). The solution was stirred under a water saturated nitrogen gas atmosphere for four hours. The peptide containing reduced cysteine was separated from the other components by chromatography over a G10  
10 Sephadex column equilibrated with 0.2 M Acetic acid. The peptide was lyophilized and stored under vacuum until used. The carrier was activated by treatment with the heterobifunctional linking agent, e.g., Epsilon-maleimidocaproic acid N-hydroxysuccinimide ester, ("EMCS"), in proportions sufficient to achieve activation of approximately 25 free amino groups per  $10^5$  molecular weight of carrier. In the  
15 specific instance of diphtheria toxoid, this amounted to the addition of 6.18 mg of EMCS (purity 75%) to each 20 mg of diphtheria toxoid.

Activation of diphtheria toxoid was accomplished by dissolving each 20 mg aliquot of diphtheria toxoid in 1 ml of 0.2 M Sodium Phosphate Buffer, pH 6.45. Aliquots of 6.18 mg EMCS were dissolved into 0.2 ml of Dimethyl Formamide  
20 ("DMF"). Under darkened conditions, the EMCS was added dropwise in 50 microliter ("ul") amounts to the DT with stirring. After 2 hours of incubation in darkness, the mixture was chromatographed on a G50 Sephadex column equilibrated with 0.1 M Sodium Citrate buffer, pH 6.0, containing 0.1 mM EDTA.

Fractions containing the EMCS activated diphtheria toxoid were concentrated  
25 over a PM 10 ultrafiltration membrane under conditions of darkness. The protein content of the concentrate was determined by either the Lowry or Bradford methods. The EMCS content of the carrier was determined by incubation of the activated carrier with cysteine-HCl followed by reaction with 10 mM of Elman's Reagent 5,5'-dithio-bis (2-nitrobenzoic acid) 10 mM. The optical density difference between a blank tube  
30 containing cysteine-HCl and the sample tube containing cysteine-HCl and carrier was translated into EMCS group content by using the molar extinction coefficient of  $13.6 \times 10^3$  for 5-thio-2-nitro benzoic acid at 412 nm.



The reduced cysteine content (-SH) of the peptide was also determined utilizing Elman's Reagent. Approximately 1 mg of peptide was dissolved in 1 ml of nitrogen gas saturated water and a 0.1 ml aliquot of this solution was reacted with Elman's Reagent. Utilizing the molar extinction coefficient of 5-thio-2-nitro-benzoic acid ( $13.6 \times 10^3$ ), the free cysteine -SH was calculated. An amount of peptide containing sufficient free -SH to react with each of the 25 EMCs activated amino groups on the carrier was dissolved in 0.1 M Sodium Citrate Buffer, pH 6.0, containing 0.1 mM EDTA, and added dropwise to the EMCS activated carrier under darkened conditions. After all the peptide solution had been added to the carrier, the mixture was incubated overnight in the dark under a water saturated nitrogen gas atmosphere.

The conjugate of the peptide linked to the carrier via EMCS is separated from other components of the mixture by chromatography over a G50 Sephadex column equilibrated with 0.2 M Ammonium Bicarbonate. The conjugate eluted in the column void volume is lyophilized and stored desiccated at 20°C until used.

The conjugate may be characterized as to peptide content by a number of methods known to those skilled in the art including weight gain, amino acid analysis, etc. Conjugates of these peptides and diphtheria toxoid produced by these methods were determined to have 20-25 moles of peptide per  $10^5$  MW of carrier and all were considered suitable as immunogens for immunization of test animals.

#### Example 4

Peptide hG17(1-9)-Ser9 was prepared by standard solid state synthesis methods. That peptide contains an amino terminal immunomimic of hG17 followed by a carboxy terminal spacer. This peptide comprises a 9 amino acid immunomimic of hG17 (pGlu-Gly-Pro-Trp-Leu-Glu-Glu-Glu-Glu-) followed by the "Ser" spacer (-Ser-Ser-Pro-Pro-Pro-Pro-Cys) attached to amino acid number 9 of the hG17 immunomimic.

The peptide was conjugated to amino groups present on the Diphtheria Toxoid ("DT") immunogenic carrier via the terminal peptide cysteine residue utilizing heterobifunctional linking agents containing a succinimidyl ester at one end and maleimide at the other end of the linking agent essentially as described in Example 4.

The immunogenic constructs of this invention include an aminoterminal (1-9) G17 peptide or an aminoterminal (1-6) G34 peptide conjugated via a peptide spacer to an immunogenic carrier. The preferred G17 sequence is pyro-Glu-Gly-Pro-Trp-Leu-Glu-Glu-Glu-Glu and the preferred G34 sequence is pGlu-Leu-Gly-Pro-Gln-Gly-Arg-Pro-Pro-Pro-Pro-Cys. The preferred spacer in both constructs is a Ser-peptide (Ser-Ser-Pro-Pro-Pro-Pro-Cys). The preferred immunogenic carrier is diphtheria toxoid, tetanus toxoid, keylimpet hemocyanin, and bovine serum albumin (BSA). The gastrin immunogen is defined as a conjugate of the pGlu- Gly-Pro-Trp-Leu-Glu-Glu-Glu-Glu peptide sequence, with an amino acid spacer linked to an immunogenic carrier. The preferred gastrin immunogen is defined as a conjugate of the (1-9) amino terminal (pGlu-Gly-Pro-Trp-Leu-Glu-Glu-Glu-Glu) peptide which is linked by peptide spacer to diphtheria toxoid.

#### Example 5

These experiments demonstrate that the immunogen induces antisera that bind to amidated G17 and glycine-extended G17, but not to G34. Specifically, this experiment demonstrates the gastrin-specificity of an antiserum raised by anti-G17 immunization of rabbits.

Antisera were absorbed onto a solid phase at a concentration of 100 µg/ml and displacement was determined in a competitive assay with a fixed concentration of radiolabelled G17 (100 pg/ml) and increasing concentrations of unlabelled ligands (1-25,000 pg/ml).

Figs. 11 and 12 show the displacement of [<sup>125</sup>I]G17 from rabbit anti-human G17 antiserum by G17, G17-Gly and G34. The antiserum used in the test depicted in Fig. 11 was obtained from animals immunized with G17(1-9):DT and was specific for the N-terminal end of G17; the antiserum for Fig. 12 was specific for the C-terminal end of G17. G17 displaced radiolabelled G17 from both antisera preparations with a 50% inhibitory concentration (IC<sub>50</sub>) of 3500 pg/ml for the rabbit anti-human G17(1-9):DT (N-terminal) and 800 pg/ml for the rabbit anti-G17 (C-terminal). Glycine-extended G17 did not displace radiolabelled G17 from the C-terminal specific antiserum, but did from the N-terminal specific antiserum (IC<sub>25</sub> 12,000 pg/ml), demonstrating that the glycine-extended G17 binds to N-terminal specific antiserum,

5           This invention and its preferred embodiments have been described in detail. One skilled in the art, upon consideration of this disclosure, may make modifications and improvements within the scope of this invention.

WE CLAIM:

1. A method for treating and/or preventing hypergastrinemia comprising administering to a patient in need thereof a therapeutically effective amount of an antigastrin immunogenic composition, comprising a G17 or G34 peptide fragment or a combination thereof linked by an amino acid spacer to an immunogenic carrier.
2. The method of claim 1, wherein the spacer is a Ser peptide spacer.
3. The method of claim 1, wherein the immunogenic carrier is selected from the group consisting of diphtheria toxoid, tetanus toxoid, and keylimpet hemocyanin.
4. The method of claim 1, wherein the hypergastrinemia is associated with pernicious anemia or administration of a substance which results in increased gastrin levels.
5. A method for treating and/or preventing hypergastrinemia comprising administering to a patient in need thereof a therapeutically effective amount of anti-G17 antibodies.
6. The method of claim 5, wherein the hypergastrinemia is associated with pernicious anemia or administration of a substance which results in increased gastrin levels.
7. A method for treating tumors associated with hypergastrinemia comprising administering to a gastrin related tumor bearing patient an anti-gastrin immunogen or anti-gastrin antibodies.
8. Use of a therapeutically effective amount of an antigastrin immunogenic composition, comprising a G17 or G34 peptide fragment or a combination thereof linked by an amino acid spacer to an immunogenic carrier in the

preparation of a medicament for treating and/or preventing hypergastrinemia in a patient.

9. Use according to claim 8, wherein the spacer is a Ser peptide spacer.

10. Use according to claim 8, wherein the immunogenic carrier is selected from the group consisting of diphtheria toxoid, tetanus toxoid, and keylimpet hemocyanin.

11. Use according to claim 8, wherein the hypergastrinemia is associated with pernicious anemia or administration of a substance which results in increased gastrin levels.

12. Use of a therapeutically effective amount of anti-G17 antibodies in the preparation of a medicament for treating and/or preventing hypergastrinemia in a patient.

13. Use according to claim 12, wherein the hypergastrinemia is associated with pernicious anemia or administration of a substance which results in increased gastrin levels.

14. Use of an anti-gastrin immunogen or anti-gastrin antibodies in the preparation of a medicament for treating tumors associated with hypergastrinemia in a gastrin related tumor bearing patient.

**Fig. 1**

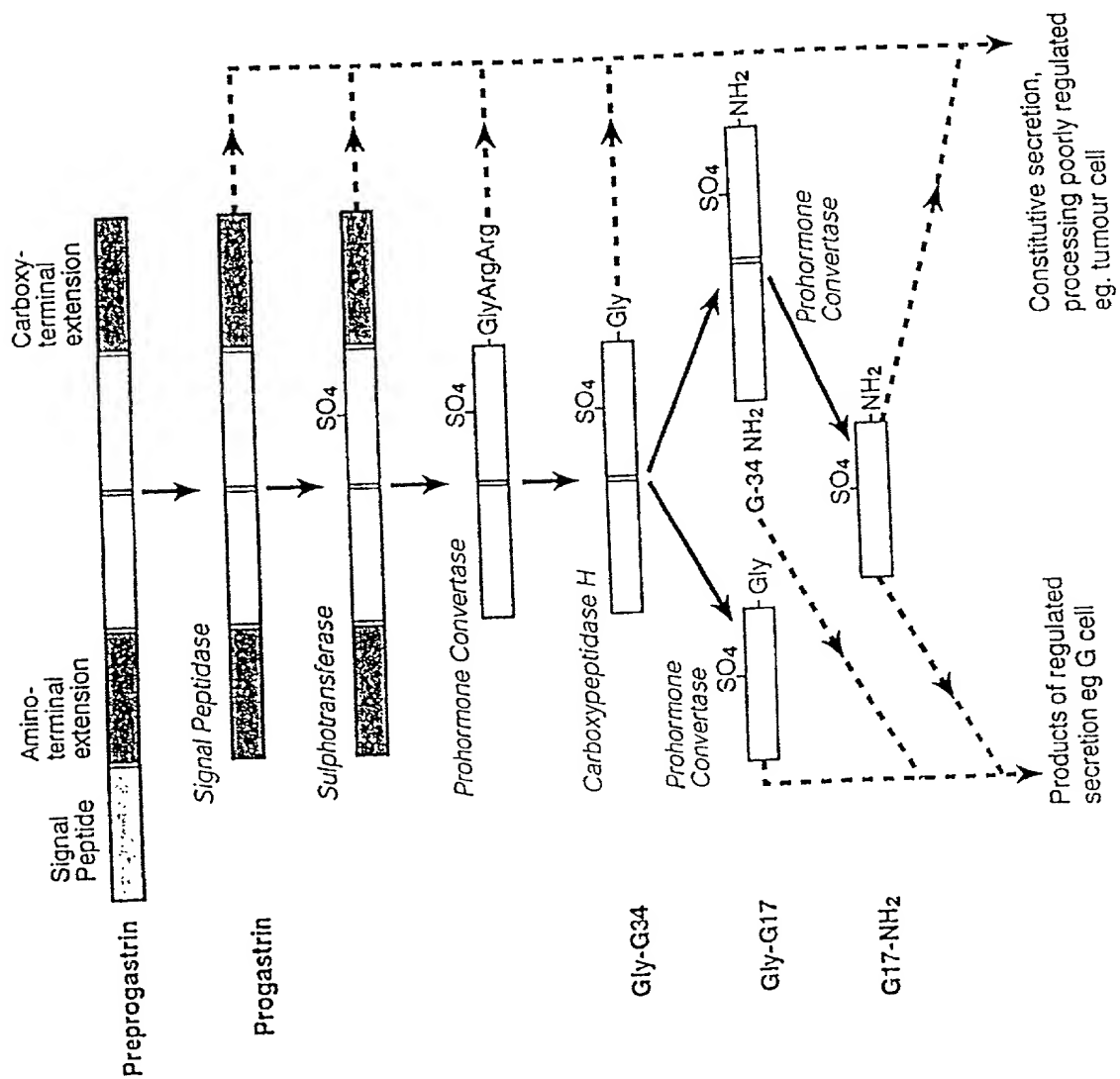
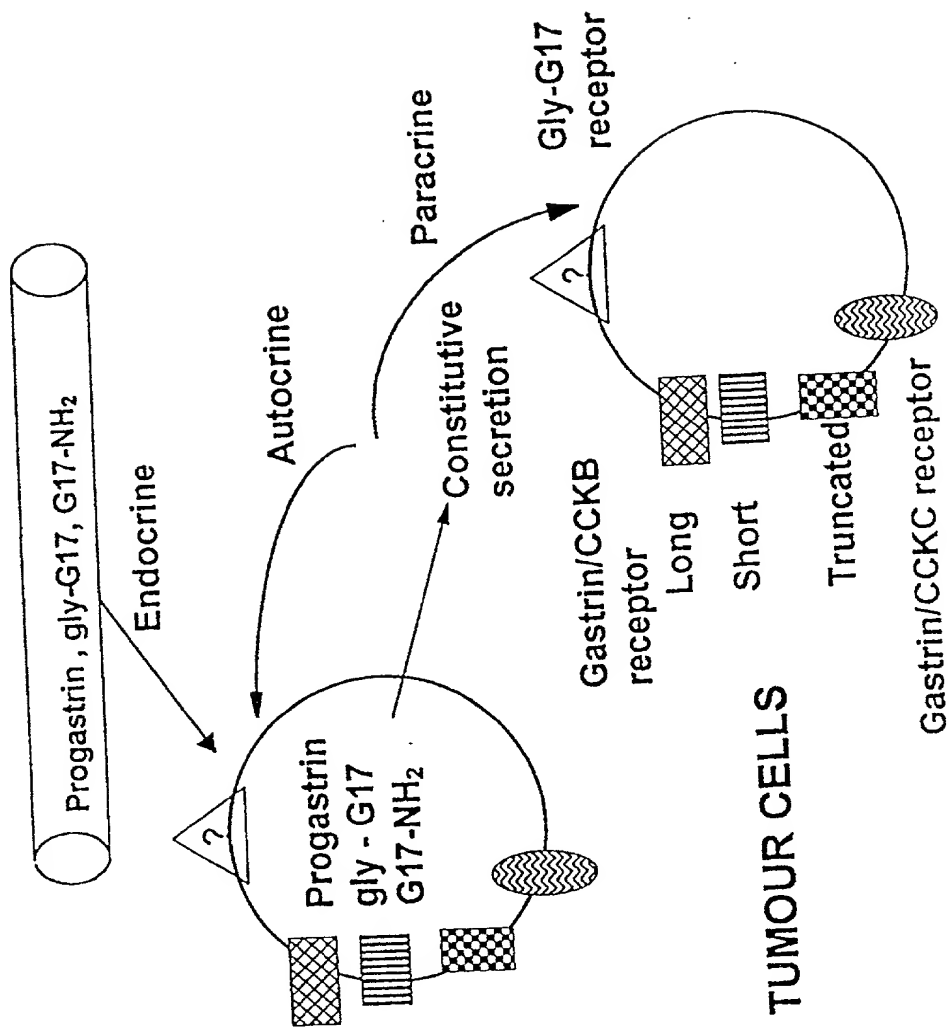
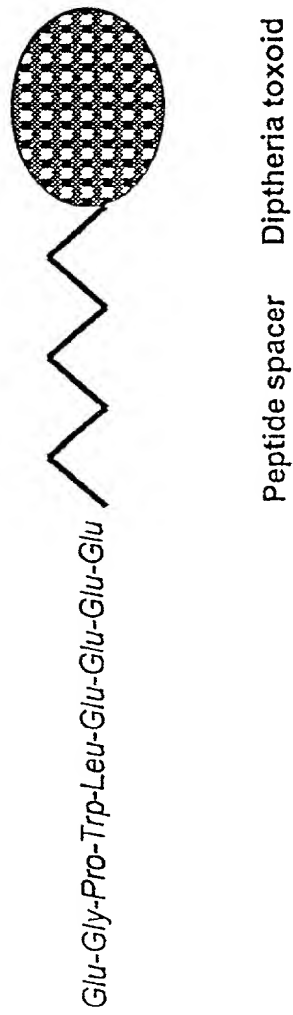


Fig. 2

**Fig. 3**





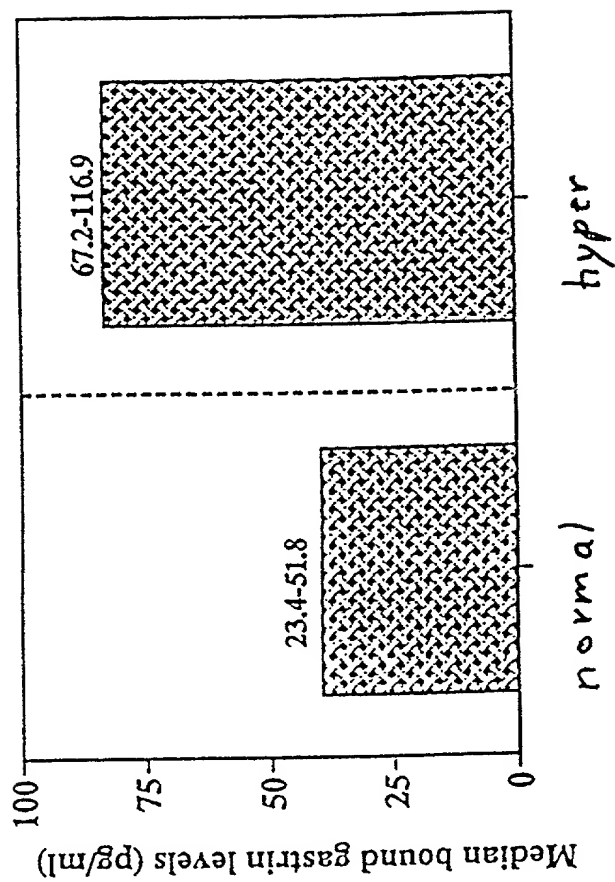
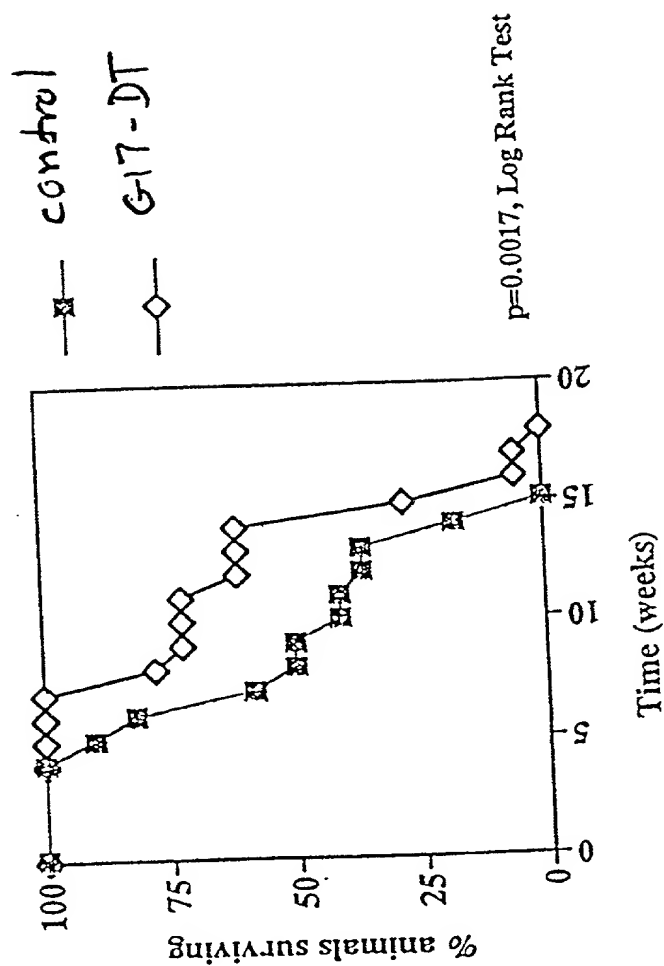


Fig. 4

Fig. 5



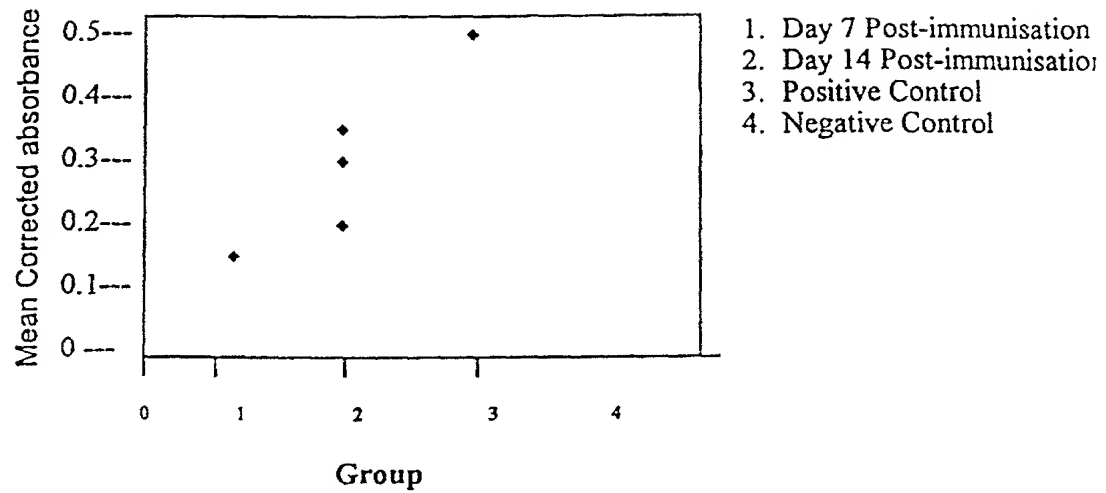
Group	Small metastases		Large metastases	
	Crypts	Tumour	Crypts	Tumour
Vehicle	6.45 (1.9)	33.3 (6.6)	6.05 (2.7)	43.2 (10.5)
G17-DT	5.87 (0.8)	26.7 (5.1)	5.80 (2.5)	25.6 (7.6)
		p<0.005		p<0.0000

[illegible]

**Fig. 7**

7/12

Timed anti - G17(1-9):DT antibody in Min mice



**Fig. 8**

8/12

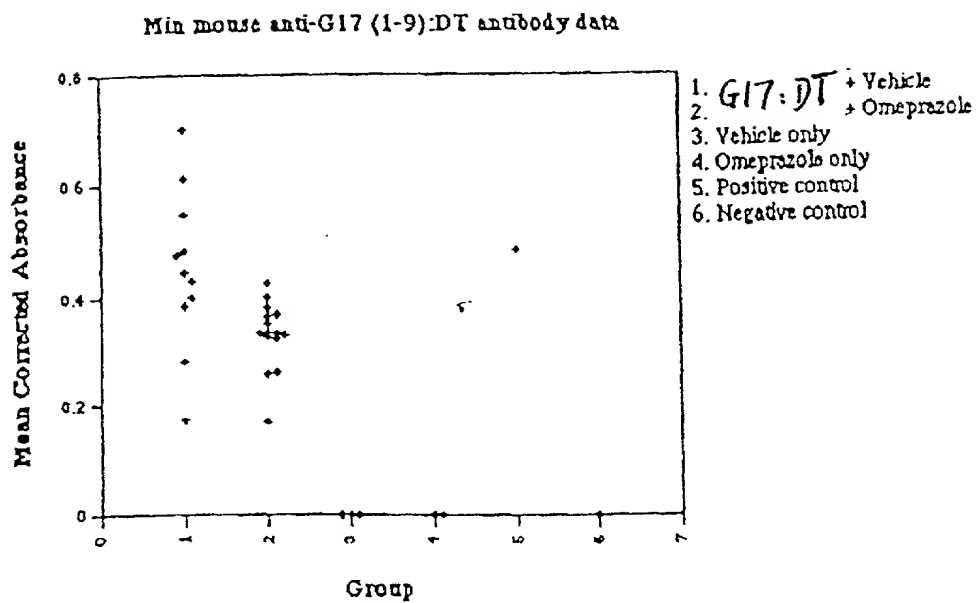
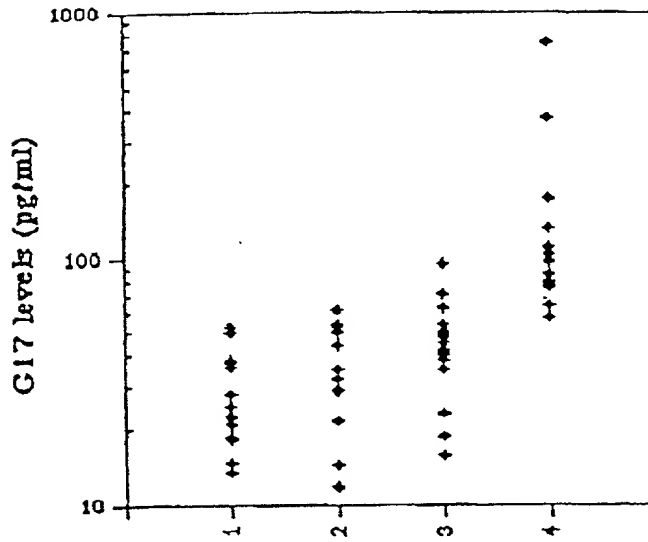
Group 1 versus Group 2,  $p=0.02$ , Mann Whitney

Fig. 9

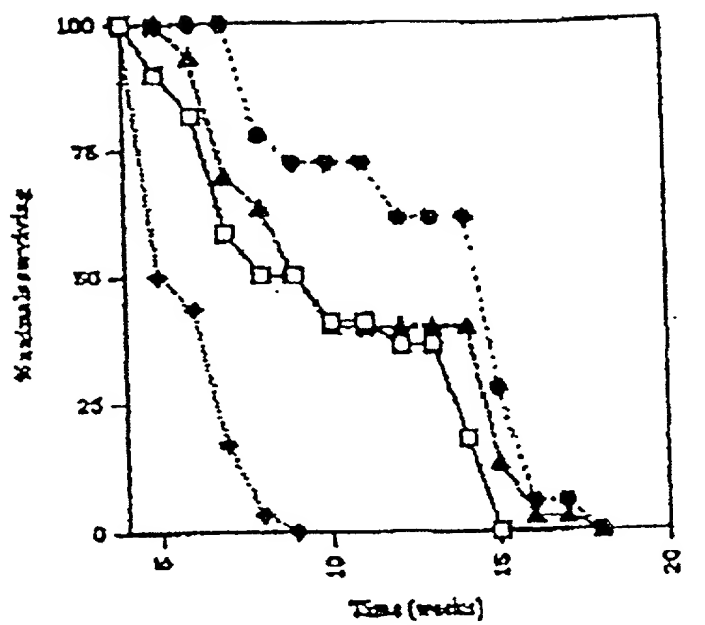
Min mouse serum G17 data



1. } hG17+DT { + Vehicle FREE  
2. } + Vehicle BOUND  
3. } + Omeprazole FREE  
4. } + Omeprazole BOUND

**Fig. 10**

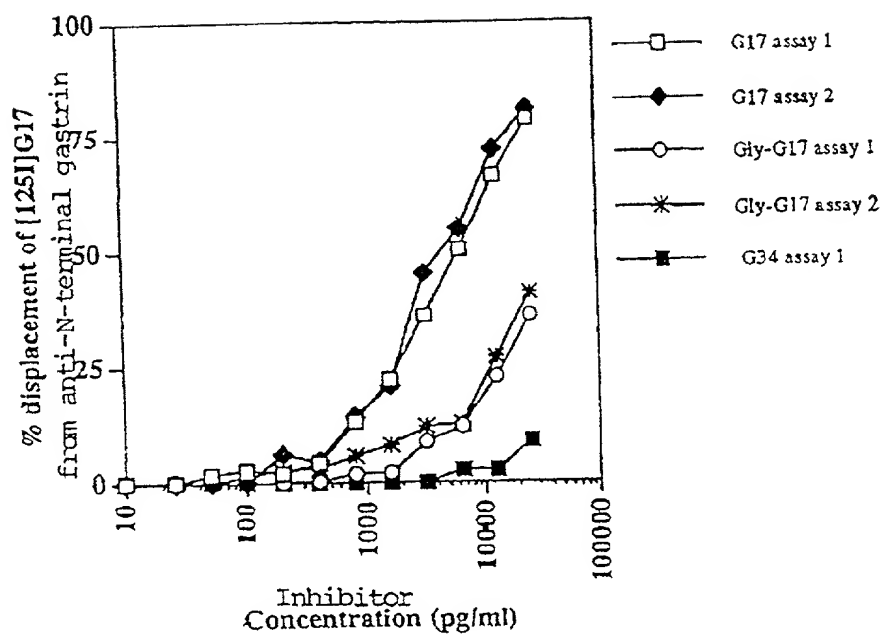
10/12



- Vehicle (oral) + Insulinogen constituents n=22  
-◆- Omeprazole (oral) + Insulinogen constituents n=20  
...◆... Vehicle (oral) + Gastrin/insulin n=18  
—▲— Omeprazole (oral) + Gastrin/insulin n=20

11/12

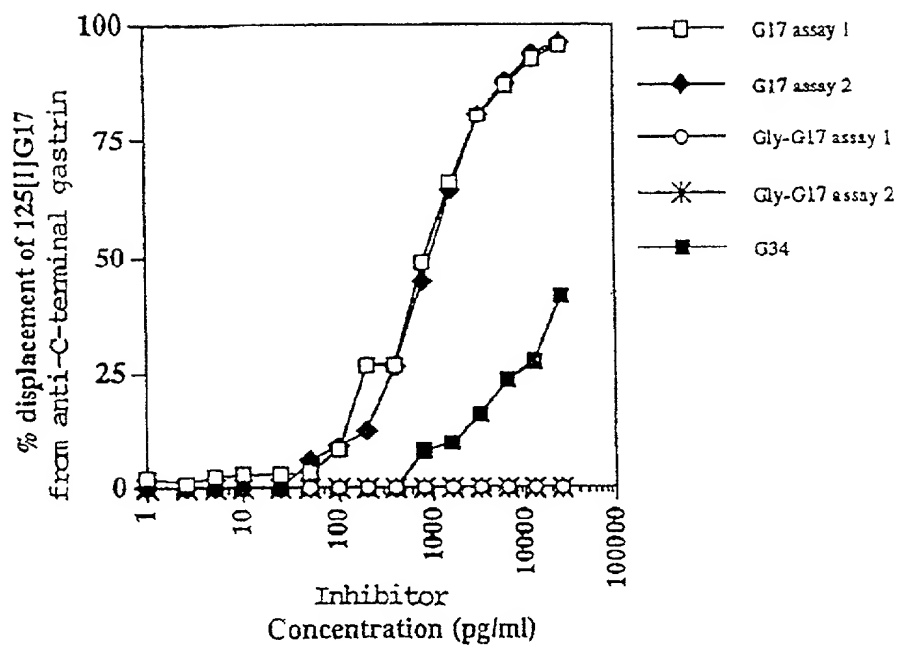
FIG. 11





12/12

FIG. 12



WO 99/59631

SEQUENCE LISTING

09/700329 }  
PCT/US99/10751  
529 Rec'd PCT/PTC 14 NOV 2000

<110> APHTON CORPORATION

<120> Prevention and Treatment of Hypergastrinemia

<130> 1102865-0035

<140> US

<141> 1999-05-14

<150> US 60/085,714

<151> 1999-05-15

<160> 4

<170> PatentIn Ver. 2.0

<210> 1

<211> 9

<212> PRT

<213> human or synthetic

<220>

<221> MOD\_RES

<222> (1)

<223> pyroglutamic acid

<400> 1

Glu Gly Pro Trp Leu Glu Glu Glu Glu

1

5

<210> 2

<211> 12

<212> PRT

<213> human or synthetic peptide

<220>

<221> MOD\_RES

<222> (1)

<223> pyroglutamic acid

<400> 2

Glu Leu Gly Pro Gln Gly Arg Pro Pro Pro Pro Cys

1

5

10

<210> 3

<211> 7

<212> PRT

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence:Synthetic  
peptide

<400> 3

Ser Ser Pro Pro Pro Cys  
1 5

<210> 4

<211> 7

<212> PRT

<213> human or synthetic peptide

<220>

<221> MOD\_RES

<222> (7)

<223> AMIDATION

<400> 4

Glu Ala Tyr Gly Trp Met Asp  
1 5

I hereby appoint the following attorneys to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith: STANLEY B. KITA, Registration No. 24,561; GEORGE A. SMITH, JR., Registration No. 24,442; WILSON OBERDORFER, Registration No. 17,379; MARY E. BAK, Registration No. 31,215, CATHY A. KODROFF, Registration Number 33,980, HENRY HANSEN, Registration No. 19,612, and WILLIAM BAK, Registration Number 37,277.

Address all telephone calls to Mary E. Bak at telephone no. (215) 540-9206. Address all correspondence to HOWSON AND HOWSON, Spring House Corporate Center, P. O. Box 457, Spring House, Pennsylvania 19477.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C. 1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full name of first inventor: Philip C. Gevas

Inventor's signature \_\_\_\_\_ Date \_\_\_\_\_

Residence: Key Biscayne, Florida 33149

Citizenship: United States of America

Post Office Address: 881 Ocean Drive #23D, Key Biscayne, Florida 33149

Full name of second inventor: Stephen Grimes

Inventor's signature \_\_\_\_\_ Date \_\_\_\_\_

Residence: Davis, California 95616

Citizenship: United States of America

Post Office Address: 551 Rutgers Drive, Davis, California 95616

Full name of third inventor: Stephen Karr

Inventor's signature \_\_\_\_\_ Date \_\_\_\_\_

Residence: Davis, California 95616

Citizenship: United States of America

Post Office Address: 2265 Halsey Circle, Davis, California 95616

Full name of fourth inventor: Dov Michaeli

Inventor's signature \_\_\_\_\_

\_\_\_\_\_  
Date

Residence: Larkspur, California 94939

Citizenship: United States of America

Post Office Address: 21 Marina Vista Avenue, Larkspur, California 94939

Full name of fifth inventor: Susan Watson

Inventor's signature \_\_\_\_\_

\_\_\_\_\_  
Date

Residence: Edwalton, Nottingham NG2 6RB Great Britain

Citizenship: United Kingdom

Post Office Address: 5 Seatolla Close, Edwalton, Nottingham NG2 6RB Great Britain

Full name of fourth inventor: Dov Michaeli

Inventor's signature \_\_\_\_\_

\_\_\_\_\_  
Date

Residence: Larkspur, California 94939

Citizenship: United States of America

Post Office Address: 21 Marina Vista Avenue, Larkspur, California 94939

Full name of fifth inventor: Susan Watson

Inventor's signature \_\_\_\_\_

\_\_\_\_\_  
Date

Residence: Edwalton, Nottingham NG2 6RB Great Britain

Citizenship: United Kingdom

Post Office Address: 5 Seatolla Close, Edwalton, Nottingham NG2 6RB Great Britain

Full name of fourth inventor: Dov Michaeli

Inventor's signature \_\_\_\_\_

\_\_\_\_\_  
Date

Residence: Larkspur, California 94939

Citizenship: United States of America

Post Office Address: 21 Marina Vista Avenue, Larkspur, California 94939

Full name of fifth inventor: Susan Watson

Inventor's signature \_\_\_\_\_

\_\_\_\_\_  
Date

Residence: Edwalton, Nottingham NG2 6RB Great Britain

Citizenship: United Kingdom

Post Office Address: 5 Seatolla Close, Edwalton, Nottingham NG2 6RB Great Britain

I hereby appoint the following attorneys to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith: STANLEY B. KITA, Registration No. 24,561; GEORGE A. SMITH, JR., Registration No. 24,442; WILSON OBERDORFER, Registration No. 17,379; MARY E. BAK, Registration No. 31,215, CATHY A. KODROFF, Registration Number 33,980, HENRY HANSEN, Registration No. 19,612, and WILLIAM BAK, Registration Number 37,277.

Address all telephone calls to Mary E. Bak at telephone no. (215) 540-9206. Address all correspondence to HOWSON AND HOWSON, Spring House Corporate Center, P. O. Box 457, Spring House, Pennsylvania 19477.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C. 1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full name of first inventor: Philip C. Gevas

Inventor's signature \_\_\_\_\_ Date \_\_\_\_\_

Residence: Key Biscayne, Florida 33149

Citizenship: United States of America

Post Office Address: 881 Ocean Drive #23D, Key Biscayne, Florida 33149

Full name of second inventor: Stephen Grimes

Inventor's signature \_\_\_\_\_ Date \_\_\_\_\_

Residence: Davis, California 95616

Citizenship: United States of America

Post Office Address: 551 Rutgers Drive, Davis, California 95616

Full name of third inventor: Stephen Karr

Inventor's signature \_\_\_\_\_ Date \_\_\_\_\_

Residence: Davis, California 95616

Citizenship: United States of America

Post Office Address: 2265 Halsey Circle, Davis, California 95616



DECLARATION AND POWER OF ATTORNEY  
FOR PATENT APPLICATION

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name,

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled PREVENTION AND TREATMENT OF HYPERGASTRINEMIA, the specification of which

(check one)                      is attached hereto  
   X        was filed on 11/14/2000 as Application No. 09/700,329 and was amended on 11/14/2000. (if applicable)

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR 1.56, including for continuation-in-part applications, material information which became available between the filing date of the prior application and the national or PCT international filing date of the continuation-in-part application.

I hereby claim foreign priority benefits under 35 U.S.C. 119(a)-(d) or 365(b) of any foreign application(s) for patent or inventor's certificate, or 365(a) of any PCT international application which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or any PCT international application having a filing date before that of the application on which priority is claimed.

Prior Foreign Application(s)			Priority Not Claimed	Certified Copy Attached?	
(Number)	(Country)	(MM/DD/YYYY)		Yes	No

I hereby claim the benefit under 35 U.S.C. 119(e) of any United States provisional application(s) listed below.

60/085,714  
(Application Number)

05/15/1998  
(Filing Date, MM/DD/YYYY)

DECLARATION AND POWER OF ATTORNEY  
FOR PATENT APPLICATION

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name,

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled PREVENTION AND TREATMENT OF HYPERGASTRINEMIA, the specification of which

(check one)                      is attached hereto  
   X        was filed on 11/14/2000 as Application No. 09/700,329 and was amended on 11/14/2000. (if applicable)

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR 1.56, including for continuation-in-part applications, material information which became available between the filing date of the prior application and the national or PCT international filing date of the continuation-in-part application.

I hereby claim foreign priority benefits under 35 U.S.C. 119(a)-(d) or 365(b) of any foreign application(s) for patent or inventor's certificate, or 365(a) of any PCT international application which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or any PCT international application having a filing date before that of the application on which priority is claimed.

Prior Foreign Application(s)			Priority Not Claimed	Certified Copy Attached?	
(Number)	(Country)	(MM/DD/YYYY)	_____	Yes	No

I hereby claim the benefit under 35 U.S.C. 119(e) of any United States provisional application(s) listed below.

60/085,714  
(Application Number)

05/15/1998  
(Filing Date, MM/DD/YYYY)

**DECLARATION AND POWER OF ATTORNEY  
FOR PATENT APPLICATION**

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name,

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled PREVENTION AND TREATMENT OF HYPERGASTRINEMIA, the specification of which

(check one)                      is attached hereto  
   X        was filed on 11/14/2000 as Application No. 09/700,329 and was amended on 11/14/2000. (if applicable)

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR 1.56, including for continuation-in-part applications, material information which became available between the filing date of the prior application and the national or PCT international filing date of the continuation-in-part application.

I hereby claim foreign priority benefits under 35 U.S.C. 119(a)-(d) or 365(b) of any foreign application(s) for patent or inventor's certificate, or 365(a) of any PCT international application which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or any PCT international application having a filing date before that of the application on which priority is claimed.

Prior Foreign Application(s)			Priority Not Claimed	Certified Copy Attached?	
(Number)	(Country)	(MM/DD/YYYY)		Yes	No

I hereby claim the benefit under 35 U.S.C. 119(e) of any United States provisional application(s) listed below.

<u>60/085,714</u>	<u>05/15/1998</u>
(Application Number)	(Filing Date, MM/DD/YYYY)

DECLARATION AND POWER OF ATTORNEY  
FOR PATENT APPLICATION

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name,

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled PREVENTION AND TREATMENT OF HYPERGASTRINEMIA, the specification of which

(check one)                      is attached hereto  
X                      was filed on 11/14/2000 as Application No. 09/700,329 and was amended on 11/14/2000. (if applicable)

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR 1.56, including for continuation-in-part applications, material information which became available between the filing date of the prior application and the national or PCT international filing date of the continuation-in-part application.

I hereby claim foreign priority benefits under 35 U.S.C. 119(a)-(d) or 365(b) of any foreign application(s) for patent or inventor's certificate, or 365(a) of any PCT international application which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or any PCT international application having a filing date before that of the application on which priority is claimed.

Prior Foreign Application(s)			Priority Not Claimed	Certified Copy Attached?	
(Number)	(Country)	(MM/DD/YYYY)		Yes	No

I hereby claim the benefit under 35 U.S.C. 119(e) of any United States provisional application(s) listed below.

<u>60/085,714</u>	<u>05/15/1998</u>
(Application Number)	(Filing Date, MM/DD/YYYY)

DECLARATION AND POWER OF ATTORNEY  
FOR PATENT APPLICATION

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name,

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled PREVENTION AND TREATMENT OF HYPERGASTRINEMIA, the specification of which

(check one)                      is attached hereto  
X                      was filed on 11/14/2000 as Application No. 09/700,329 and was amended on 11/14/2000. (if applicable)

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR 1.56, including for continuation-in-part applications, material information which became available between the filing date of the prior application and the national or PCT international filing date of the continuation-in-part application.

I hereby claim foreign priority benefits under 35 U.S.C. 119(a)-(d) or 365(b) of any foreign application(s) for patent or inventor's certificate, or 365(a) of any PCT international application which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or any PCT international application having a filing date before that of the application on which priority is claimed.

Prior Foreign Application(s)			Priority Not Claimed	Certified Copy Attached?	
(Number)	(Country)	(MM/DD/YYYY)		Yes	No

I hereby claim the benefit under 35 U.S.C. 119(e) of any United States provisional application(s) listed below.

<u>60/085,714</u>	<u>05/15/1998</u>
(Application Number)	(Filing Date, MM/DD/YYYY)

I hereby appoint the following attorneys to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith: STANLEY B. KITA, Registration No. 24,561; GEORGE A. SMITH, JR., Registration No. 24,442; WILSON OBERDORFER, Registration No. 17,379; MARY E. BAK, Registration No. 31,215; CATHY A. KODROFF, Registration Number 33,980; HENRY HANSEN, Registration No. 19,612, and WILLIAM BAK, Registration Number 37,277.

Address all telephone calls to Mary E. Bak at telephone no. (215) 540-9206. Address all correspondence to HOWSON AND HOWSON, Spring House Corporate Center, P. O. Box 457, Spring House, Pennsylvania 19477.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C. 1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full name of first inventor: Philip C. Gevas

Inventor's signature  12-16-00  
Date

Residence: Key Biscayne, Florida 33149

Citizenship: United States of America

Post Office Address: 881 Ocean Drive #23D, Key Biscayne, Florida 33149

Full name of second inventor: Stephen Grimes

Inventor's signature \_\_\_\_\_  
Date

Residence: Davis, California 95616

Citizenship: United States of America

Post Office Address: 551 Rutgers Drive, Davis, California 95616

Full name of third inventor: Stephen Karr

Inventor's signature \_\_\_\_\_  
Date

Residence: Davis, California 95616

Citizenship: United States of America

Post Office Address: 2265 Halsey Circle, Davis, California 95616

I hereby appoint the following attorneys to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith: STANLEY B. KITA, Registration No. 24,561; GEORGE A. SMITH, JR., Registration No. 24,442; WILSON OBERDORFER, Registration No. 17,379; MARY E. BAK, Registration No. 31,215, CATHY A. KODROFF, Registration Number 33,980, HENRY HANSEN, Registration No. 19,612, and WILLIAM BAK, Registration Number 37,277.

Address all telephone calls to Mary E. Bak at telephone no. (215) 540-9206. Address all correspondence to HOWSON AND HOWSON, Spring House Corporate Center, P. O. Box 457, Spring House, Pennsylvania 19477.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C. 1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full name of first inventor: Philip C. Gevas

Inventor's signature \_\_\_\_\_ Date \_\_\_\_\_

Residence: Key Biscayne, Florida 33149

Citizenship: United States of America

Post Office Address: 881 Ocean Drive #23D, Key Biscayne, Florida 33149

Full name of second inventor: Stephen Grimes

Inventor's signature *Stephen Grimes* 18 Dec, 2000  
Date

Residence: Davis, California 95616

Citizenship: United States of America

Post Office Address: 551 Rutgers Drive, Davis, California 95616

Full name of third inventor: Stephen Karr CA

Inventor's signature \_\_\_\_\_ Date \_\_\_\_\_

Residence: Davis, California 95616

Citizenship: United States of America

Post Office Address: 2265 Halsey Circle, Davis, California 95616

I hereby appoint the following attorneys to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith: STANLEY B. KITA, Registration No. 24,561; GEORGE A. SMITH, JR., Registration No. 24,442; WILSON OBERDORFER, Registration No. 17,379; MARY E. BAK, Registration No. 31,215, CATHY A. KODROFF, Registration Number 33,980, HENRY HANSEN, Registration No. 19,612, and WILLIAM BAK, Registration Number 37,277.

Address all telephone calls to Mary E. Bak at telephone no. (215) 540-9206. Address all correspondence to HOWSON AND HOWSON, Spring House Corporate Center, P. O. Box 457, Spring House, Pennsylvania 19477.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C. 1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full name of first inventor: Philip C. Gevas

Inventor's signature \_\_\_\_\_ Date \_\_\_\_\_

Residence: Key Biscayne, Florida 33149

Citizenship: United States of America

Post Office Address: 881 Ocean Drive #23D, Key Biscayne, Florida 33149

Full name of second inventor: Stephen Grimes

Inventor's signature \_\_\_\_\_ Date \_\_\_\_\_

Residence: Davis, California 95616

Citizenship: United States of America

Post Office Address: 551 Rutgers Drive, Davis, California 95616

Full name of third inventor: Stephen Karr

Inventor's signature Stephen Karr 25 January 2001  
Date

Residence: Davis, California 95616 CH

Citizenship: United States of America

Post Office Address: 2265 Halsey Circle, Davis, California 95616



400

Full name of fourth inventor: Dov Michaeli

Inventor's signature *Dov Michaeli* 1/8/01  
Date

Residence: Larkspur, California 94939

Citizenship: United States of America

Post Office Address: 21 Marina Vista Avenue, Larkspur, California 94939

Full name of fifth inventor: Susan Watson

Inventor's signature \_\_\_\_\_  
Date

Residence: Edwalton, Nottingham NG2 6RB Great Britain

Citizenship: United Kingdom

Post Office Address: 5 Seatolla Close, Edwalton, Nottingham NG2 6RB Great Britain

Patented by the United States Patent and Trademark Office

Full name of fourth inventor: Dov Michaeli

Inventor's signature \_\_\_\_\_ Date \_\_\_\_\_

Residence: Larkspur, California 94939

Citizenship: United States of America

Post Office Address: 21 Marina Vista Avenue, Larkspur, California 94939

5700 Full name of fifth inventor: Susan Watson

Inventor's signature S. A. Watson <sup>GB</sup> 18th Dec 2000.  
Date

Residence: Edwalton, Nottingham NG2 6RB Great Britain

Citizenship: United Kingdom

Post Office Address: 5 Seatolla Close, Edwalton, Nottingham NG2 6RB Great Britain